

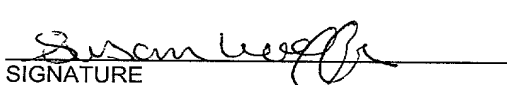


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FORM PTO-1390 (REV 5-93)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE		ATTORNEY'S DOCKET NO. 05032.86955	
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371				U.S. APPLICATION NO. (If known, See 37 CFR 1.5) TBA 09/508852	
INTERNATIONAL APPLICATION NO. PCT/NL98/00634		INTERNATIONAL FILING DATE 3 November 1998		PRIORITY DATE CLAIMED 3 November 1997	
TITLE OF INVENTION REMOVABLE PROTECTIVE COATING					
APPLICANT(S) FOR DO/EO/US Antoon J. G. van ROSSUM et al.					
Applicant herewith submits to the United State Designated/Elected Office (DO/EO/US) the following items and other information:					
<ol style="list-style-type: none"><li>1. <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. 371.</li><li>2. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371.</li><li>3. <input checked="" type="checkbox"/> This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).</li><li>4. <input checked="" type="checkbox"/> A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.</li><li>5. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371(c)(2))<ol style="list-style-type: none"><li>a. <input type="checkbox"/> is transmitted herewith (required on if not transmitted by the International Bureau).</li><li>b. <input checked="" type="checkbox"/> has been transmitted by the International Bureau.</li><li>c. <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US).</li></ol></li><li>6. <input type="checkbox"/> A translation of the International Application into English (35 U.S.C. 371(c)(2)).</li><li>7. <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))<ol style="list-style-type: none"><li>a. <input type="checkbox"/> are transmitted herewith (required only if not transmitted by the International Bureau).</li><li>b. <input type="checkbox"/> have been transmitted by the International Bureau.</li><li>c. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired.</li><li>d. <input checked="" type="checkbox"/> have not been made and will not be made.</li></ol></li><li>8. <input type="checkbox"/> A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).</li><li>9. <input type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).</li><li>10. <input type="checkbox"/> A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).</li></ol>					
Items 11-16 below concern other document(s) or information included:					
<ol style="list-style-type: none"><li>11. <input checked="" type="checkbox"/> An Information Disclosure Statement under 37 C.F.R. 1.97 and 1.98.</li><li>12. <input type="checkbox"/> An Assignment document for recording. A separate cover sheet in compliance with 37 C.F.R. 3.28 and 3.31 is included.</li><li>13. <input checked="" type="checkbox"/> A FIRST preliminary amendment. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment.</li><li>14. <input type="checkbox"/> A substitute specification.</li><li>15. <input type="checkbox"/> A change of power of attorney and/or address letter.</li><li>16. <input checked="" type="checkbox"/> Other items or information:  International Search Report (ISA/EPO)</li></ol>					

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U.S. APPLICATION NO. (If known, See 37 CFR 1.5) TBA 09/508852				INTERNATIONAL APPLICATION NO PCT/NL98/00634		ATTORNEY'S DOCKET NO. 05032.86955		
17. ■ The following fees are submitted:  <b>Basic National Fee (37 CFR 1.492(a)(1)-(5):</b> Search Report has been prepared by the EPO or JPO \$840.00 International preliminary examination fee paid to USPTO (37 CFR 1.482) \$670.00 No International preliminary examination fee paid to USPTO (37 CFR 1.482), but international search fee paid to USPTO (37 CFR 1.445(a)(2)) \$760.00 Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO \$970.00 International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(1)-(4) \$ 96.00					CALCULATIONS		PTO USE ONLY	
ENTER APPROPRIATE BASIC FEE AMOUNT =					\$840.00			
Surcharge of \$130.00 for furnishing the oath or declaration later than 20 or 30 months from the earliest claimed priority date (37 CFR 1.492(e)).					\$0.00			
CLAIMS		NUMBER FILED		NUMBER EXTRA		RATE		
Total Claims		27 -20 =		7		X \$ 18.00		
Independent Claims		2 - 3 =		0		X \$ 78.00		
Multiple dependent claims (if applicable)				\$260.00		\$0.00		
TOTAL OF ABOVE CALCULATIONS =					\$966.00			
Reduction by 1/2 for filing by small entity, if applicable. Verified Small Entity statement must also be filed (note 37 CFR 1.9, 1.27, 1.28).					\$0.00			
SUBTOTAL =					\$966.00			
Processing fee of \$130.00 for furnishing the English translation later than 20 or 30 months from the earliest claimed priority date (37 CFR 1.492(f)).					\$0.00			
TOTAL NATIONAL FEE =					\$966.00			
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property.					\$0.00			
TOTAL FEES ENCLOSED =					\$966.00			
					Amount to be:			
					refunded		\$	
					charged		\$	
a. ■ A check in the amount of \$ <u>966.00</u> to cover the above fees is enclosed. b. □ Please charge my Deposit Account No. 19-0733 in the amount of \$_____ to cover the above fees. A duplicate copy of this sheet is enclosed. c. ■ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 19-0733. A duplicate copy of this sheet is enclosed.								
NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.								
SEND ALL CORRESPONDENCE TO:					 SIGNATURE			
Banner & Witcoff, Ltd. Eleventh Floor 1001 G Street, N.W. Washington, D.C. 20001-4597					Susan A. Wolfe Registration No. 33,568			
Telephone: (202) 508-9100					March 17, 2000			

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**IN THE UNITED STATES PATENT AND TRADEMARK OFFICE**

In re the Application of

Antoon J. G. van ROSSUM et al.

Serial No.: TBA

Filed: Herewith (March 17, 2000)

For: REMOVABLE PROTECTIVE COATING

Atty. Docket: 05032.86955

U.S. National Stage  
International Application No.:  
PCT/NL98/00634

**PRELIMINARY AMENDMENT**

Assistant Commissioner of Patents  
Washington, D. C. 20231

**BOX PCT**

Sir:

Prior to calculation of claim fees and examination, please amend the application as follows:

**IN THE ABSTRACT:**

Please insert the following Abstract at the end of the specification, which is also provided herewith on a separate sheet:

**--ABSTRACT OF THE DISCLOSURE**

A protective coating which is removable with a removing agent comprising a strong base, which protective coating comprises a pigment, a weak base and a binder, the binder being a polymer having a weight-average molecular weight of 10,000-100,000 and an acid value of 40-250. A protective agent and a method for forming the protective coating.--

**IN THE SPECIFICATION:**

Page 1, between the title of the application and the first paragraph, insert the heading --Background of the Invention--.

Page 3, between lines 16 and 17, insert the heading --Brief Description of the Invention--.

Page 4, between lines 15 and 16, insert the heading --Detailed Description of the Invention--.

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Page 12, between lines 5 and 6, insert the heading --Examples--.

**IN THE CLAIMS:**

Claim 3, line 1, delete “or 2”.

Claim 4, lines 1 and 2, delete “any one of the preceding claims” and insert therefor --claim 1--.

Claim 5, lines 1 and 2, delete “any one of the preceding claims” and insert therefor --claim 1--.

Claim 6, lines 1 and 2, delete “any one of the preceding claims” and insert therefor --claim 1--.

Claim 8, lines 1 and 2, delete “any one of the preceding claims” and insert therefor --claim 1--.

Claim 9, lines 1 and 2, delete “any one of the preceding claims” and insert therefor --claim 1--.

Claim 10, lines 1 and 2, delete “any one of the preceding claims” and insert therefor --claim 1--.

Claim 11, lines 1 and 2, delete “any one of the preceding claims” and insert therefor --claim 1--.

Claim 13, lines 1 and 2, delete “any one of the preceding claims” and insert therefor --claim 1--.

Claim 14, lines 1 and 2, delete “any one of the preceding claims” and insert therefor --claim 1--.

Claim 15, lines 5 and 6, delete “any one of the preceding claims” and insert therefor --claim 1--.

Claim 18, line 4, delete “claims 15-17” and insert therefor --claim 15--.

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Claim 20, line 2, delete "claims 1-14" and insert therefor --claim 15--.

Claim 22, line 1, delete "or 21".

Claim 23, line 1, delete "claims 20-22" and insert therefor --claim 20--.

Claim 26, line 2, delete "claims 15-17" and insert therefor --claim 15--; and line 3, delete "claims 20-25" and insert therefor --claim 20--.

Cancel claims 27 and 28 and insert new claim 29, as follows:

-- 29. A method of removing a protective agent from a transparent surface comprising applying benzyl alcohol to said surface.--

**REMARKS**

By this amendment, the specification is amended to place the application in preferred form; the claims are amended to remove multiple dependencies; and an Abstract of the Disclosure is provided as required under 37 C.F.R. § 1.72(b). Examination on the merits of the instant application is respectfully requested.

Respectfully submitted,



Susan A. Wolffe  
Reg. No. 33,568

Date: March 17, 2000

BANNER & WITCOFF, LTD.  
Eleventh Floor  
1001 G Street, N.W.  
Washington, D. C. 20001-4597  
(202) 508-9100

Attachment: Abstract of the Disclosure

SAW:lab

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### ABSTRACT OF THE DISCLOSURE

A protective coating which is removable with a removing agent comprising a strong base, which protective coating comprises a pigment, a weak base and a binder, the binder being a polymer having a weight-average molecular weight of 10,000-100,000 and an acid value of 40-250. A protective agent and a method for forming the protective coating.

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**VERIFIED STATEMENT (DECLARATION) CLAIMING SMALL ENTITY STATUS**  
**(37 CFR 1.9(f) and 1.27(c)) — SMALL BUSINESS CONCERN**

Applicant or Patentee: Raadgevend Chemiebureau RSB v.o.f.

Serial No.: \_\_\_\_\_ Filing Date: \_\_\_\_\_

Patent No.: \_\_\_\_\_ Issued: \_\_\_\_\_

For: \_\_\_\_\_

**"Removable protective coating"**

I hereby declare that I am

- ☐ the owner of the small business concern identified below:  
☒ an official of the small business concern empowered to act on behalf of the concern identified below:

NAME OF CONCERN A.J.G. van Rossum A. BertelsADDRESS OF CONCERN Geerstraat 8a, 5111 PS BAARLE-NASSAU

I hereby declare that the above identified small business concern qualifies as a small business concern as defined in 13 CFR 121.3-18, and reproduced in 37 CFR 1.9(d), for purposes of paying reduced fees under section 41(a) and (b) of Title 35, United States Code, in that the number of employees of the concern, including those of its affiliates, does not exceed 500 persons. For purposes of this statement, (1) the number of employees of the business concern is the average over the previous fiscal year of the concern of the persons employed on a full-time, part-time or temporary basis during each of the pay periods of the fiscal year, and (2) concerns are affiliates of each other when either, directly or indirectly, one concern controls or has the power to control the other, or a third party or parties controls or has the power to control both.

I hereby declare that rights under contract or law have been conveyed to and remain with the small business concern identified above with regard to the invention, entitled "Removable protective coating" by inventor(s)

described in

- ☐ the application filed herewith  
☒ application serial no. \_\_\_\_\_, filed March 17, 2000  
☐ patent no. \_\_\_\_\_, issued \_\_\_\_\_

If the rights held by the above identified small business concern are not exclusive, each individual, concern or organization having rights to the invention is listed below\* and no rights to the invention are held by any person, other than the inventor, who could not qualify as a small business concern under 37 CFR 1.9(d) or by any concern which would not qualify as a small business concern under 37 CFR 1.9(d) or a nonprofit organization under 37 CFR 1.9(e).

\*NOTE: Separate verified statements are required from each named person, concern or organization having rights to the invention averring to their status as small entities. (37 CFR 1.27)

NAME \_\_\_\_\_

ADDRESS \_\_\_\_\_

☐ INDIVIDUAL ☐ SMALL BUSINESS CONCERN ☐ NONPROFIT ORGANIZATION

NAME \_\_\_\_\_

ADDRESS \_\_\_\_\_

☐ INDIVIDUAL ☐ SMALL BUSINESS CONCERN ☐ NONPROFIT ORGANIZATION

I acknowledge the duty to file, in this application or patent, notification of any change in status resulting in loss of entitlement to small entity status prior to paying, or at the time of paying, the earliest of the issue fee or any maintenance fee due after the date on which status as a small entity is no longer appropriate. (37 CFR 1.28(b))

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application, any patent issuing thereon, or any patent to which this verified statement is directed.

NAME OF PERSON SIGNING A.J.G. van Rossum A. BertelsTITLE OF PERSON OTHER THAN OWNER partnerADDRESS OF PERSON SIGNING Geerstraat 8a 5111 PS BAARLE-NASSAU

SIGNATURE \_\_\_\_\_

DATE March 3, 2000A. RossumA. Bertels

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Title: Removable protective coating

This invention relates to a removable protective coating, and to a protective agent and a method for forming such coating.

In horticulture, many plants are grown in  
5 greenhouses. Thus, they can benefit from optimized  
conditions, such as temperature, amount of light, humidity  
and the like. However, a problem here is that the plants  
being grown are exposed, under warm, sunny weather  
conditions, to a large amount of radiation, which might  
10 disturb the living conditions and growth of the plants, and  
might even cause them to burn. To obviate this problem, it  
is customary in horticulture to protect the plants during  
the spring and the summer against the adverse effect of  
undue radiation by providing the transparent surfaces of  
15 the greenhouse with a protective coating.

One of the chief requirements to be met by such a  
protective coating is that sufficient protection from light  
and heat is achieved. To meet this requirement, the  
protective coating should contain a pigment, such as chalk  
20 or titanium oxide. A further important requirement is that  
the protective agent from which the protective coating is  
formed exhibits sufficient adhesion to the surface of a  
greenhouse. When its adhesive strength is too low, the  
coating will not be resistant to weather influences and it  
25 will be necessary to restore the coating several times per  
season or to replace it. When adhesion is too strong, it  
requires much effort to remove the coating at the end of  
the season.

In European patent application 0 428 937, it is  
30 proposed to use a protective agent based on a polymeric  
binder and pigments, such as aluminum pigments, titanium  
pigments or calcium carbonate. No particular requirements  
are set with regard to the nature of the polymeric binder.

In practice, it has been found that many polymeric  
35 binders impart too high an adhesive strength to the

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protective agent. This makes it particularly labor-intensive to remove the protective coating at the end of the season and/or requires the use of aggressive chemicals, which is undesirable on the ground of health and environmental considerations.

European patent application 0 478 067 discloses a method for applying a protective coating to a greenhouse, which coating provides protection against solar radiation. A thin layer of a protective agent is applied to a surface, and after a chemical conversion, a stable coating is obtained. To remove the coating at the end of the season, the coating is treated with a stripping agent, which removes the stability and the water-resistance of the coating. The surface can subsequently be easily cleaned by rinsing with water.

The protective agent used according to this European patent application is based on an inorganic pigment and a polymeric binder. The single example shows the use of calcium carbonate as pigment and a partially esterified copolymer of styrene and maleic acid anhydride as binder.

A protective coating formed from the agent described in EP-A-0 478 067 proves to be insufficiently resistant to weather influences. Such a coating must be frequently restored or even be re-applied several times in the course of one season. It is assumed that the instability of the coating is due to degradation of the polymer chains of the binder by, for instance, UV radiation. This degradation shortens the chains of the binder, thereby reducing the adhesive strength of the protective coating. The adverse effect of degradation of the polymer chains of the binder is all the greater in the above-mentioned copolymer of styrene and maleic acid anhydride because the chains of that polymer are already short as it is.

European patent application 0 533 367 discloses a coating based on a copolymer of an  $\alpha,\beta$ -ethylenically unsaturated carboxylic acid, another  $\alpha,\beta$ -ethylenically

unsaturated monomer and an amphiphilic monomer. In the examples of the application, the coating described was tested on steel plates. The properties that are deemed to be relevant are drying time, resistance to acid rain and to  
5 rust of metallic powders, and removability by an aqueous, basic solution.

European patent application 0 578 498 relates to a specific acrylic polymer of improved heat resistance and processability. The polymer has a number-average molecular  
10 weight of 1,000 to 1,000,000, a glass transition temperature of -80°C or higher, and a molecular weight distribution of 5 or less. The intended uses of the polymer are *inter alia* acrylic rubbers, pressure-sensitive adhesives, dispersants, additives to asphalt, coatings,  
15 fiber processing agents, and improvers for resins and water inks.

It is an object of the present invention to provide a protective coating which affords protection against (solar) radiation and has an adhesive strength such that  
20 the above-described disadvantages from the prior art do not occur. What is further contemplated is that the protective coating can be easily removed at the moment when it is no longer desired.

It has been found that these objects are achieved  
25 when a protective coating is formed from a protective agent which comprises a pigment and a specific binder. Accordingly, the invention relates to a protective coating which is removable with a removing agent comprising a strong base and a complex former, which protective coating  
30 comprises a pigment and a binder, the binder being a polymer having a weight-average molecular weight of 10,000-100,000 and an acid value of 40-250.

Surprisingly, a protective coating according the invention has been found to have excellent adhesive power.  
35 The coating is sufficiently stable to afford prolonged protection against radiation, without requiring interim

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repair. Further, the coating is eminently resistant to various weather influences, such as rain, frost and UV radiation. Another great advantage of a protective coating according to the invention is that it can be removed in a manner that is not particularly labor-intensive nor requires chemicals that affect the environment unacceptably and/or constitute health hazards.

The protective coating according to the invention can be applied to surfaces of different materials. Preferably, the surface is a substantially transparent surface, such as an outside surface of a greenhouse, for instance a horticultural greenhouse. Typically, the surface will be made of glass or plastic. Conventionally used plastics are, for instance, polycarbonates, polyolefins, polyethylene terephthalate and polyesters.

As stated, the protective coating according to the invention comprises a pigment and a binder. Preferably, the pigment is present in an amount of 30-95% by weight, more preferably of 40-85% by weight, based on the weight of the protective coating. The binder is preferably present in an amount of 4-60% by weight, more preferably of 6-45% by weight, based on the weight of the protective coating. It has been found that a protective coating containing these components in these amounts has both optimum protective action and optimum adhesive strength.

As pigment, any substance can be used that can be suitably dispersed in a protective agent from which the protective coating according to the invention is formed, and which imparts to the coating the desired protective action against (solar) radiation. Preferably, a pigment is used which yields a white protective coating. Suitable pigments are selected from the group of calcium carbonate, titanium oxide, a silicate, such as magnesium or aluminum silicate, gypsum, baryte, and combinations thereof. Depending on the desired properties of the protective coating, the skilled person will be able to suitably select

the pigment. Titanium oxide has a very high covering power, so that only a relatively small quantity thereof is needed. As a consequence, the ratio between binder and pigment is more favorable. When much binder with respect to the pigment can be used, the weather resistance of the protective coating is better. The advantage of the use of calcium carbonate is that a protective coating based thereon becomes slightly transparent in damp weather, so that the light intensity within a greenhouse adjusts itself to the weather conditions. Further, calcium carbonate is an economically attractive natural product and upon removal gives little, if any, visual or other contamination of the environment.

An important constituent of the protective coating according to the invention is the polymeric binder. This constituent should be selected such that a sufficient adhesion of the coating to a surface is obtained, while yet the coating can be easily removed at any desired time. According to the invention, as binder, a polymer is used which has a weight-average molecular weight of 10,000-100,000 and an acid value of 40-250.

The weight-average molecular weight of the binder should not be too low. A low weight-average molecular weight is normally indicative of short polymer chains, which will entail a poorer binding effect in the protective agent to be used. Moreover, the degradation of the binder under the influence of (UV) radiation has more effect when short chains are broken than when long chains are broken. Too high a weight-average molecular weight also entails disadvantages. Often, the viscosity of the protective agent will be too high when the chains of the binder are too long. The agent is then difficult to apply to a surface. It is preferred that the weight-average molecular weight of the polymeric binder lies between 15,000, preferably 20,000, and 75,000, preferably 50,000. The weight-average molecular weight can be suitably determined with gel

permeation chromatography (GPC). This technique, which is known per se, can be carried out in this connection in a manner known to those skilled in the art, using, for instance, tetrahydrofuran as solvent and polystyrene as  
5 external standard.

Another requirement to be satisfied by the binder according to the invention is that it have an acid value of 40 to 250. The acid value is connected with the average number of free acid groups per chain of the polymeric  
10 binder. This value is preferably 60-160. The acid value can be determined by titration with potassium hydroxide, with the end point being determined potentiometrically. The acid value then corresponds to the number of milligrams of potassium hydroxide used in the titration per gram of the  
15 polymeric binder.

A polymer which, according to the invention, is suitable for use as binder in a protective agent preferably has a polydispersity of a value of 2 to 6, preferably of 3 to 5. The term 'polydispersity' used herein means the ratio  
20 between the weight-average and the number-average molecular weight ( $M_w/M_n$ ). The number-average molecular weight, like the weight-average molecular weight, can be determined using GPC.

Preferably, the binder has a glass transition  
25 temperature between 10 and 60°C, with a particular preference for the range between 20 and 50°C. It has been found that a protective coating based on a polymer satisfying this requirement exhibits particularly suitable adhesion to a surface of a greenhouse. It has also been  
30 found that using, as a binder, a polymer having a glass transition temperature within the range mentioned yields a protective agent that has good handling properties and is easy to apply to form the protective coating.

As binder, both homopolymers and copolymers can be  
35 used. The nature of the monomers in the binder is of less importance than the parameters already mentioned.

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The binder is preferably a vinyl polymer. The use of a vinyl polymer as binder yields an exceptionally suitable protective coating. Suitable monomers are, for instance, vinyl aromatic monomers, such as  $\alpha$ -methylstyrene and

5 styrene, acrylonitrile, methacrylonitrile, acrylamide, vinyl acetate, vinyl chloride, phenoxyethyl acrylate, multifunctional acrylates, such as hexanediol dimethyl acrylate, glycol dimethyl acrylate, divinyl benzene and esters of methacrylic acid or acrylic acid, or mixtures of

10 these esters. Examples of suitable esters comprise alkyl esters, where the alkyl group can contain from 1 to 20 carbon atoms, alkoxy alkyl esters, such as butoxy ethyl acrylate and butoxy alkyl methacrylate, and hydroxy alkyl esters. An acrylic acid or methacrylic acid ester can

15 account for up to 90% of the polymer. When acrylonitrile or acrylamide is present in the vinyl polymer, these monomers will typically be present in the polymer in an amount of less than 10 to 15% by weight. Styrene and vinyl acetate, respectively, may account for up to 30 and 50% by weight of

20 the polymer. Other suitable monomers are acid monomers such as acrylic acid, methacrylic acid, maleic acid, fumaric acid, crotonic acid, itaconic acid, aconitic acid and semi-esters thereof, and maleic acid anhydride and the like. These acid monomers can be present in amounts of up

25 to 50% by weight. Particularly suitable monomers are methyl methacrylate, butyl acrylate, 2-ethylhexyl acrylate, ethyl acrylate, styrene, methacrylic acid and acrylic acid.

Further, it is possible that the polymer is wholly or partly crosslinked. Eligible as crosslinking monomers

30 are methylolacrylamide, methylolmethacrylamide and the like. These crosslinking monomers will typically account for not more than 5% by weight of the polymer.

In addition to a pigment, a weak base and a binder, a protective coating according to the invention can further

35 comprise a few other components.

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Thus, it is advantageous to include an adhesion promoter in the protective coating. The adhesion of the coating to a surface will be improved by the presence of an adhesion promoter, while the ease of removing the protective coating is hardly, if at all, affected. Depending on the material of the surface on which a protective coating is to be provided, the skilled person will be able to select a suitable adhesion promoter. Preferably, the adhesion promoter must be water-soluble and contain an amino group. For use on glass surfaces, it is recommended to use a silane, such as  $\gamma$ -aminopropyltriethoxy silane,  $\gamma$ -aminopropyltrimethoxy silane,  $\gamma$ -(methylamino)propyltrimethoxy silane,  $\gamma$ -aminopropylmethyldiethoxy silane,  $\gamma$ -(2-aminoethyl-3-aminopropyl)triethoxy silane and  $\gamma$ -(2-aminoethyl-3-aminopropyl)methyldimethoxy silane. An adhesion promoter will typically be present in the protective agent in an amount of 0.05 to 1% by weight, preferably from 0.1 to 0.3% by weight, based on the weight of the protective coating.

Another constituent yielding advantages is a pigment divider. The presence of such a substance prevents aggregation of pigment upon drying of the protective agent which has been applied in diluted form for forming the protective coating. A pigment divider can be present in amounts of from 0.1 to 0.5% by weight, based on the weight of the protective coating. The nature of the pigment divider depends on the nature of the pigment present in the protective coating. Thus, sodium hexametaphosphate is highly suitable when calcium carbonate is used as pigment. When titanium dioxide is used as pigment, for instance a polymeric multifunctional surfactant, such as Ser-Ad FA 607® (available from the firm of Hüls AG) can be used excellently as a pigment divider.

To optimize the viscosity of the protective agent for forming a protective coating according to the invention, a thickener can be included. The optimum

viscosity of the protective agent depends on the method by which the coating is applied to a surface. If a surface is brushed with a protective agent, a higher viscosity will be desirable than when a surface is sprayed with a protective agent. Further, the viscosity must be sufficient to obtain a thick coating. On the basis of his common professional knowledge, the skilled person will be able to determine which viscosity is most suitable in any given case.

Examples of thickeners comprise organic and inorganic thickeners, such as hydroxyethyl cellulose, magnesium aluminum silicate and combinations thereof. The amount of the thickener will be tuned to the desired viscosity and typically be between 1 and 5% by weight, based on the weight of the protective coating.

To be mentioned as other additional constituents of the protective coating are dispersants, detergents, antifoam agents, preserving agents and the like.

It will be clear that the invention also relates to a protective agent for forming a protective coating as described above. The protective agent will contain water, in addition to the protective coating constituents already mentioned. If the protective agent contains calcium carbonate as pigment, it will contain such an amount of water that the above-mentioned constituents of the protective coating in the ratios specified, are diluted 1.5-5 times. If titanium oxide is used as pigment, that dilution is a factor of 5-12 times. It is an advantage of the invention that no volatile organic solvents need to be used.

Preferably, there is also a weak base present in the protective agent according to the invention. It provides for neutralization of free acid groups present in one or more components of the agent. It has also been found that the presence of the weak base leads to improved film formation of the protective agent upon drying, when a protective coating is being formed. Preferably, the weak



base is selected from the group of ammonia, mono- di- and trialkylamines, with the alkyl group containing from 1 to 8 carbon atoms. Particularly preferred is ammonia.

The weak base is preferably present in an amount of 5 0.2-5% by weight, more preferably of 0.4-3% by weight, based on the weight of the protective agent.

As stated, the application of the protective agent to form a protective coating can occur in different ways. Possible ways include spraying, brushing and the like. The 10 protective action of the coating will be hardly, if at all, affected by the manner of application.

When in the course of time, for instance at the end of the season, the protective coating is to be removed, the coating according to the invention is treated with a 15 removing agent, comprising a strong base and a complex former. The removing agent renders the binder in the protective coating water-soluble.

The strong base is preferably present in the removing agent in an amount of from 2 to 10% by weight, 20 more preferably of from 2 to 5% by weight, based on the weight of the removing agent. Suitable strong bases are, for instance, alkali metal hydroxides such as sodium hydroxide, potassium hydroxide and lithium hydroxide. Preferably, sodium hydroxide is used.

25 Preferably present, in addition to the strong base, is a complex former, preferably in an amount of 2 to 10% by weight, based on the weight of the removing agent. It has been found that especially the use of the trisodium salt of nitrilo-triacetic acid or the tetrasodium salt of ethylene 30 diamine tetraacetic acid greatly promotes the protective coating becoming soluble. The use of trisodium salt of nitrilo-triacetic acid is preferred. This substance is properly and rapidly biodegradable.

Especially when a protective coating provided on a 35 plastic is to be removed, it is found to be of great advantage to use a removing agent that further comprises an

organic solvent. It is also possible to use an organic solvent separately, in addition to the removing agent. The latter option is advantageous in that the organic solvent used does not necessarily need to be alkali-resistant. A  
5 great many solvents are eligible for use as a separate component. Examples include benzyl alcohol, tetrahydrofuran, 1,4-dioxane, dimethyl sulfoxide, higher alcohols, such as butanol, pentanol, hexanol, cyclohexanol and isomers thereof, and cyclohexanone. The organic solvent  
10 effects a still easier removal of the protective coating.

It is preferred to use the organic solvent in the removing agent. This is beneficial in particular to the simplicity of the procedure of removing the protective coating. In that case, an alkali-resistant organic solvent  
15 should be used. The amount of organic solvent is preferably 10-30% by weight, more preferably 15-25% by weight, based on the weight of the removing agent. Particularly preferred is the use of benzyl alcohol. Benzyl alcohol is little volatile, little toxic to man and animals and hardly  
20 combustible, so that the health of those working with the removing agent is not put at risk. When benzyl alcohol ends up in the environment after the removal of the protective coating, this does not yield unacceptable contamination.

In addition to the constituents mentioned, the  
25 removing agent may further contain a thickener, such as xanthan gum. Xanthan gum renders the removing agent highly pseudoplastic, so that it is thin when being applied and thick after being applied. This property prevents the agent from flowing off the surface too fast. Further, the  
30 removing agent may contain a substance reducing the surface tension, or an emulsifier. For instance, the sodium salt of dodecylbenzenesulfonic acid is suitable.

To remove the protective coating, the coating is treated with the above-described removing agent. This  
35 treatment comprises suitably applying, as by spraying or pouring, the removing agent onto the coating to be removed.

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Typically, the removing agent is used in a 5 to 10-fold dilution. After application of the removing agent, the surface can be rinsed with water. It is also possible to allow the rain to wash things off. Thereafter, virtually all traces of the protective coating will be gone.

The invention will presently be further elucidated in and by the following examples.

Example 1: Preparation of a protective agent

In a reaction vessel, the following substances were added to each other in the order specified:

	wt. %
water	36.9
alkyl ethoxylate (non-ionic surfactant)	0.2
antifoam agent	0.3
30% solution of sodium hexametaphosphate	0.6
ammonia solution 25%	1.5
polymer dispersion *)	14.8
finely divided calcium carbonate	44

The mixture obtained was stirred for 20 minutes under high shear conditions.

Then the following substances were added, in succession:

magnesium aluminum silicate	0.8
hydroxyethyl cellulose	0.6
preserving agent	0.1
60% solution of $\gamma$ -aminopropyltriethoxy silane	0.2

Again, vigorous stirring was carried out, until a homogeneous product was formed.

\*) The polymer dispersion used had a solids content of 45% and an acid value of 73, calculated on the solids content.

The polymer in the binder had been formed exclusively from acrylate monomers.

20 Pa.sec (Brookfield 0.5 rpm, spindle 5)

5 20 mPa.sec (Brookfield 0.5 rpm, spindle 1)

In a reaction vessel, the following substances were added  
10 to each other in the order specified:

15 The mixture obtained was stirred at a moderate speed, until  
a thickening had occurred.

20	sodium hydroxide solution 33%	6.0
	solution of trisodium nitrilotriacetate 40%	20
	30% solution of an amphoteric surfactant	0.5

Viscosity of end product 20°C:

30 Pa.sec (Brookfield 0.5 rpm, spindle 5)

30 Viscosity of dilution of use at 20°C (1 part by weight of product diluted with five parts by weight of water):

40 mPa.sec (Brookfield 0.5 rpm, spindle 1)

The protective agent of Example 1 was diluted with 1.5 parts by weight of water and applied to a glass horticultural greenhouse by spraying in the early spring.

After six months, the protective coating formed was still fully intact.

Example 4: Removing a protective coating

- 5 The removing agent of Example 2 was diluted, for use, with five parts by weight of water and applied to the protective coating of Example 3 by spraying in the autumn. After a few showers of rain, which had come down by the end of a few days, the protective coating was found to have been  
10 completely washed off by the rain.

Example 5: Preparation of a protective agent

In a reaction vessel, the following substances were added to each other in the order specified:

15		wt. %
	water	6.0
	Ser-AD FA 607® (Hüls AG)	0.3
	alkyl ethoxylate (non-ionic surfactant)	0.3
	polymer dispersion *)	3.0
20	antifoam agent	0.5
	titanium white	20

The mixture obtained was dispersed with a high shear dissolver until the desired fineness was achieved.

25

Then the following were added, in succession:

	water	31.3
	hydroxyethyl cellulose	0.90

- 30 Stirring was performed until a homogeneous mixture was obtained. Then the following were added, in succession:
- |    |   |     |
|----|---|-----|
|    | ammonia solution 25%                                  | 3.4 |
|    | polymer dispersion *)                                 | 34  |
|    | preserving agent                                      | 0.1 |
| 35 | 60% solution of $\gamma$ -aminopropyltriethoxy silane | 0.2 |

Vigorous stirring was performed until a homogeneous product was formed.

- \*) The polymer dispersion used had a solids content of 45% and an acid value of 73, calculated on the solids content. The polymer in the binder had been formed exclusively from acrylate monomers.

Viscosity of end product at 20°C:

- 10 Pa.sec (Brookfield 0.5 rpm, spindle 5)  
Viscosity of dilution of use at 20°C (1 part by weight of product diluted with four parts by weight of water):  
15 mPa.sec (Brookfield 0.5 rpm, spindle 1)

15 Example 6: Preparation of a removing agent

In a reaction vessel, the following substances were added to each other in the order specified:

	wt. %
water	57.65
20 Xanthan gum	0.5

The mixture obtained was stirred at a moderate speed, until a thickening was achieved.

Then there were added, in succession:

25 Dodecylbenzenesulfonic acid	0.25
sodium hydroxide solution 33%	6.1
solution of trisodium nitrilotriacetate 40%	10
benzyl alcohol	25
pine oil	0.5

- 30 Stirring was performed until a homogeneous product was formed.

Viscosity of end product 20°C:

10 Pa.sec (Brookfield 0.5 rpm, spindle 5)

Viscosity of dilution of use at 20°C (1 part by weight of product diluted with five parts by weight of water):

20 mPa.sec (Brookfield 0.5 rpm, spindle 1)

5

Example 7: Forming a protective coating

The protective agent of Example 5 was diluted with four parts by weight of water and applied to a plastic horticultural tunnel by spraying, in the early spring.

10 After six months, the protective coating formed was still fully intact.

Example 8: Removing a protective coating

15 The removing agent of Example 6 was diluted with five parts by weight of water and applied by spraying onto the protective coating of Example 7, in the autumn. After a short time of action, the protective coating was removed completely with a water jet.

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## CLAIMS

1. A protective coating removable with a removing agent comprising a strong base and a complex former, which protective coating comprises a pigment and a binder, the binder being a polymer having a weight-average molecular weight of 10,000-100,000 and an acid value of 40-250.
2. A protective coating according to claim 1, wherein the binder has a weight-average molecular weight of 15,000-75,000, preferably of 20,000-50,000.
3. A protective coating according to claim 1 or 2, wherein the acid value of the binder is between 60 and 160.
4. A protective coating according to any one of the preceding claims, wherein the binder has a polydispersity of 2-6.
5. A protective coating according to any one of the preceding claims, wherein the glass transition temperature of the binder is between 10 and 60°C, preferably between 20 and 50°C.
6. A protective coating according to any one of the preceding claims, wherein the binder is a vinyl polymer.
7. A protective coating according to claim 6, wherein the vinyl polymer is based on one or more monomers selected from the group of methyl methacrylate, butyl acrylate, 2-ethylhexyl acrylate, ethyl acrylate, styrene, methacrylic acid, and acrylic acid.
8. A protective coating according to any one of the preceding claims, wherein the binder is present in an amount of 4-60% by weight, based on the weight of the protective coating.
9. A protective coating according to any one of the preceding claims, wherein the pigment is selected from the group of calcium carbonate, titanium oxide, a silicate, such as magnesium or aluminum silicate, gypsum, baryte, and combinations thereof.



10. A protective coating according to any one of the preceding claims, wherein the pigment is present in an amount of 30-95% by weight, based on the weight of the protective coating.
- 5 11. A protective coating according any one of the preceding claims, wherein the protective coating further comprises an adhesion promoter.
12. A protective coating according to claim 11, wherein the adhesion promoter is selected from the group of
- 10 silanes.
13. A protective coating according to any one of the preceding claims, wherein the protective coating further comprises a pigment divider.
14. A protective coating according to any one of the
- 15 preceding claims, wherein the protective coating further comprises a thickener.
15. A protective agent comprising a pigment and a water-carried binder, wherein the binder is a polymer having a weight-average molecular weight of 10,000-100,000
- 20 and an acid value of 40-250, for forming a removable protective coating according to any one of the preceding claims.
16. A protective agent according to claim 15, wherein the protective agent further comprises a weak base selected
- 25 from the group of ammonia, mono-, di- and trialkylamines, with the alkyl group containing from 1 to 8 carbon atoms.
17. A protective agent according to claim 16, wherein the weak base is present in an amount of 0.2-5% by weight, based on the weight of the protective agent.
- 30 18. A method for forming a protective coating which is removable with a removing agent comprising a strong base and a complex former, wherein a protective agent according to claims 15-17 is applied to a substantially transparent surface, which protective agent, after drying, forms the
- 35 protective coating.

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19. A method according to claim 18, wherein the substantially transparent surface is an outside wall surface of a greenhouse.

20. A method for removing a protective coating according to claims 1-14, wherein the protective coating is treated with a removing agent which comprises a strong base and a complex former.

21. A method according to claim 20, wherein the strong base is present in an amount of 1-10% by weight, based on the weight of the removing agent.

22. A method according to claim 20 or 21, wherein the complex former is present in an amount of 2-10% by weight, based on the weight of the removing agent.

23. A method according to claims 20-22, wherein the removing agent further comprises an organic solvent.

24. A method according to claim 23, wherein the organic solvent is selected from the group of tetrahydrofuran, benzyl alcohol, and higher alcohols.

25. A method according to claim 24, wherein the removing agent comprises 10-30% by weight, based on the weight of the removing agent, of benzyl alcohol.

26. A kit comprising a protective agent according to claims 15-17 and a removing agent, which removing agent is suitable for use in a method according to claims 20-25.

27. Use of a polymer having a weight-average molecular weight of 10,000-100,000 and an acid value of 40-250 as binder in a protective agent.

28. Use of benzyl alcohol for removing a protective agent from a transparent surface.

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## DECLARATION AND POWER OF ATTORNEY FOR U.S. PATENT APPLICATIONS

( ) Original ( ) Supplemental ( ) Substitute (x) PCT

As a below named inventor, I hereby declare that: my residence, post office address and citizenship are as stated below next to my name; that I verily believe that I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural inventors are named below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

"Removable Protective Coatings"

Title: \_\_\_\_\_

which is described and claimed in:

( ) the attached specification, or  
 (x) the specification in the application Serial No. \_\_\_\_\_ filed March 17, 2000 ;  
 and with amendments through \_\_\_\_\_ (if applicable),  
 (x) the specification in International Application No. PCT/NL98/00634 \_\_\_\_\_, filed  
 November 3, 1998 , and as amended on \_\_\_\_\_ (if applicable).

I hereby state that I have reviewed and understand the content of the above-identified specification, including the claims, as amended by any amendment(s) referred to above.

I acknowledge my duty to disclose information of which I am aware which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, §1.56(a).

I hereby claim foreign priority benefits under Title 35, United States Code, §119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

COUNTRY	APPLICATION NO.	DATE OF FILING	PRIORITY CLAIMED
the Netherlands	1007433	November 3, 1997	(x) YES ( ) NO
_____	_____	_____	( ) YES ( ) NO
_____	_____	_____	( ) YES ( ) NO
_____	_____	_____	( ) YES ( ) NO
_____	_____	_____	( ) YES ( ) NO
_____	_____	_____	( ) YES ( ) NO

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, §1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

SERIAL NO.	U.S. FILING DATE	STATUS
_____	_____	( ) Patented ( ) Pending ( ) Abandoned
_____	_____	( ) Patented ( ) Pending ( ) Abandoned
_____	_____	( ) Patented ( ) Pending ( ) Abandoned

As a named inventor I hereby appoint the following attorney(s) and/or agents to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith: P.T.O.

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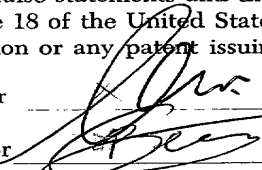
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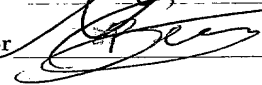
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I further declare that all statements made herein of my own knowledge are true, and that all statements on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

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